

pantothenate and vitamin B₆ content, and its label failed to bear such information concerning its vitamin properties as has been determined to be, and by regulations prescribed as, necessary in order fully to inform purchasers as to its value for such uses, since its label failed to bear the statement, "The need for calcium pantothenate and vitamin B₆ in human nutrition has not been established," as required by the regulations.

On June 1, 1943, the William T. Thompson Co., claimant, having admitted the allegations of the libel, judgment of condemnation was entered and the product was ordered released under bond for relabeling in conformance with the law, under the supervision of the Food and Drug Administration.

5779. Adulteration and misbranding of elixir thiamine hydrochloride. U. S. v. 52 Bottles of Elixir Thiamine Hydrochloride. Decree of condemnation. Product ordered delivered to charitable institutions. (F. D. C. No. 9591. Sample No. 23501-F.)

Examination showed that this product contained substantially less than 250 International Units (USP units) of vitamin B₁ per fluid ounce.

On March 19, 1943, the United States attorney for the Eastern District of Pennsylvania filed a libel against 52 bottles, each containing 1 gallon of the above-named product, at Philadelphia, Pa., alleging that the article had been shipped on or about February 2, 1943, from Newark, N. J., by the Standard Drug Co.; and charging that it was adulterated and misbranded. A portion of the article (35 bottles) was labeled in part: "Standard Elixir Vitamin B₁ N. J. F. Elixir Thiamin Hydrochloride. Each fluid ounce contains 500 Intern. Units Vitamin B₁." The remainder of the article (17 bottles) was relabeled by the consignee, and at the commencement of the libel proceedings was labeled in part: "Elixir Thiamin Hydrochloride * * * Each fluid ounce contains: Thiamine Hydrochloride—1.5 mg. (equivalent to Vitamin B-1—500 Units)."

The article was alleged to be adulterated in that a valuable constituent, vitamin B₁, had been in whole or in part omitted or abstracted therefrom.

It was alleged to be misbranded in that the following statements: (In the case of the portion bearing the original label) "Each fluid ounce contains 500 Intern. Units Vitamin B₁," and (in the case of the relabeled portion) "Each Fluid ounce Contains: Thiamine Hydrochloride—1.5 mg. (equivalent to Vitamin B-1—500 Units)," were false since the article contained a lesser amount of vitamin B₁ per fluid ounce. It was alleged to be misbranded further in that it purported to be a food for special dietary use by reason of its vitamin B₁ content, and its label failed to bear such information concerning its vitamin properties as had been determined to be, and by regulations prescribed as, necessary in order fully to inform purchasers as to its value for such uses, since its label failed to bear a statement of the proportion of the minimum daily requirement of vitamin B₁ supplied by a specified quantity of the article when consumed as directed during a period of 1 day.

The article was also alleged to be adulterated and misbranded under the provisions of the law applicable to drugs, as reported in the notices of judgment on drugs and devices.

On May 10, 1943, no claimant having appeared, judgment of condemnation was entered and the product was ordered to be delivered to charitable institutions.

5780. Adulteration and misbranding of Ocean-Lax. U. S. v. 29 Bottles of Ocean-Lax. Decree of condemnation and destruction. (F. D. C. No. 6368. Sample Nos. 40885-E, 40886-E.)

On December 6, 1941, the United States attorney for the Eastern District of Pennsylvania filed a libel against 29 bottles of Ocean-Lax at Philadelphia, Pa., alleging that the article had been shipped in interstate commerce within the period from on or about July 3 to August 11, 1941, by Mineralized Foods, Inc., from Baltimore, Md.; and charging that it was adulterated and misbranded.

The article was alleged to be adulterated in that it contained deleterious substances, the laxative drugs, senna pods, purging cassia, and rhubarb root, which might render it injurious to health.

It was alleged to be misbranded in that the following statement, appearing on the label, created the impression that the article was appropriate for food purposes, whereas, because of its content of cathartic drugs, it was not suitable for such purpose "* * * Consists of an imported rare variety of Sea Vegetables high in alkalinity and food minerals carefully blended with * * * Each Ocean-Lax Tablet averages approximately 1½ milligrams of natural organic